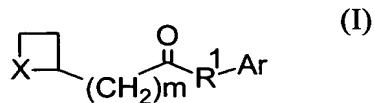


IN THE CLAIMS

1. (Original) Compound having the general formula (I):



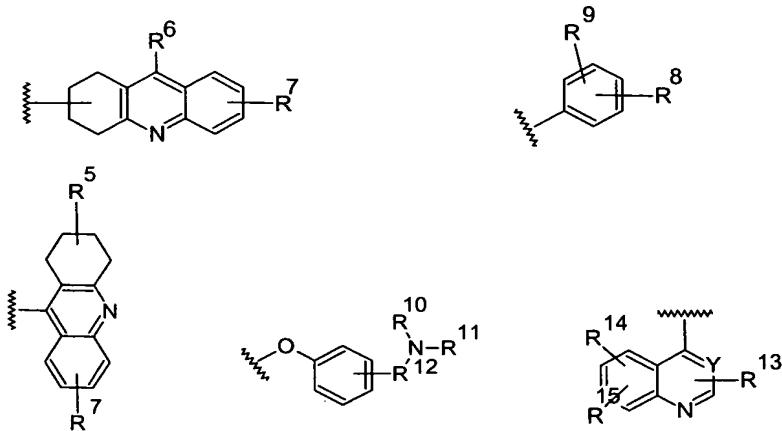
or its geometric isomers, its optically active forms, diastereoisomers, its racemic forms, or its pharmaceutically acceptable salts, wherein R^1 is selected from the group consisting of: C₂-C₉ alkandiamine, C₂-C₆ amine; X is selected from the group consisting of: -S-S-, -S-, -CH₂-, -CH₂-CH₂-; m is an integer greater than zero and lower than eight; Ar represents an aromatic group; R^1 comprises a nitrogen linked directly to the carbonyl.

2. (Original) Compound according to claim 1, wherein X represents -S-S-.

3. (Currently amended) Compound according to claim 1 or 2, wherein m is an integer greater than two and lower than five.

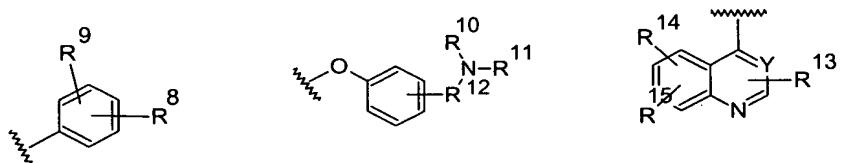
4. (Original) Compound according to claim 3, wherein m is four.

5. (Currently amended) Compound according to ~~one of the previous claims~~ claim 1, wherein Ar presents a formula selected from the group consisting of:

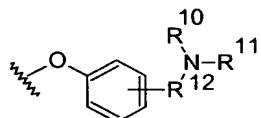


wherein R⁵ is selected from the group consisting of: hydrogen, amine, nitroalkyl, -NH₂, nitro, halogen, hydroxy; R⁶ is selected from the group consisting of: hydrogen, amine, alkandiamine, -NH₂; R⁷ is selected from the group consisting of: hydrogen, group having an electron attractor inductive effect; R¹³, R¹⁴, R¹⁵, R⁸ and R⁹ are selected, each independently of the others, from the group consisting of: hydrogen, hydroxy, halogen, alkoxy, alkyl, nitroalkyl, cyanoalkyl, nitro, cyano; R¹⁰ and R¹¹, are selected, each independently of the other, from the group consisting of: hydrogen, C₁-C₄ alkyl; R¹² represents a C₁-C₄ alkyl; Y is selected from the group consisting of -CH- and -N-.

6. (Original) Compound according to claim 5, wherein Ar presents a formula selected from the group consisting of:



7. (Original) Compound according to claim 6, wherein Ar presents the formula:

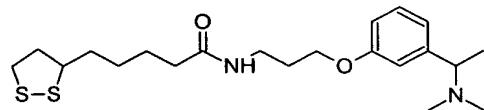


wherein R¹ represents a C₂-C₆ amine.

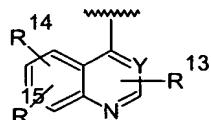
8. (Original) Compound according to claim 7, wherein R¹ presents the formula -N(CH₂)_n-⁻, wherein the nitrogen is directly linked to the carbonyl and n is an integer greater than one and smaller than five.

9. (Original) Compound according to claim 8, wherein n is three; R¹⁰ and R¹¹ represent, each, a respective methyl; R¹² represents an ethyl and is linked at the meta position with respect to the oxygen.

10. (Original) Compound according to claim 9, and having the following formula:



11. (Original) Compound according to claim 6, wherein Ar presents the formula:

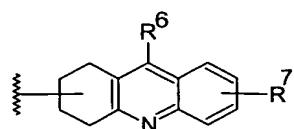


wherein Y represents N, R¹ represents an alkandiamine having the formula –NR³-R²-NR⁴-; R² represents a C₂-C₅ alkyl; R³ and R⁴ are selected, each independently of the other, from the group consisting of: hydrogen, methyl; R¹³, R¹⁴, R¹⁵ are selected, each independently of the others, from the group consisting of: hydrogen, hydroxy, halogen, C₁-C₄ alkoxy, C₁-C₄ alkyl.

12. (Original) Compound according to claim 11, wherein R² represents a linear propyl; R³ and R⁴ each represent a hydrogen; R¹³ represents a halogen; R¹⁴ and R¹⁵ are selected, each independently of the other, from the group consisting of: halogen, hydroxy, C₁-C₄ alkoxy.

13. (Currently amended) Compound according to claim 11 or 12, wherein R¹³ represents a chlorine; R¹⁴ and R¹⁵ represent, each, a respective methoxy.

14. (Original) Compound according to claim 5, wherein Ar presents the formula:

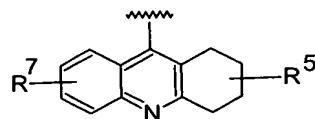


R⁷ is selected from the group consisting of: hydrogen, C₁-C₄ alkoxy, halogen; R⁶ is selected from the group consisting of: –NH₂, alkandiamine, amine; R¹ represents a C₁ amine.

15. (Original) Compound according to claim 14, wherein R⁶ is selected from the group consisting of: –NH₂ and amine C₁-C₄.

16. (Original) Compound according to claim 14, wherein R⁷ is a chlorine situated in position 6; R⁶ represents -NH₂; R¹ represents -NH-CH₂-, wherein the nitrogen is linked to the carbonylic carbon.

17. (Original) Compound according to claim 5, wherein Ar presents the formula:



wherein R¹ represents a C₂-C₆ alkandiamine.

18. (Original) Compound according to claim 17, wherein R¹ represents a C₃-C₄ alkandiamine.

19. (Currently amended) Compound according to claim 17 or 18, wherein R¹ presents the formula -NR³-R²-NR⁴-, wherein R² represents a C₂-C₄ alkyl, R³ and R⁴ are selected, each independently of the other, from the group consisting of: hydrogen, methyl.

20. (Original) Compound according to claim 19, wherein R³ and R⁴ represent, each, a respective hydrogen.

21. (Currently amended) Compound according to claim 19 or 20, wherein R² represents -(CH₂)₃-.

22. (Currently amended) Compound according to ~~one of the claims from~~ ~~claim 17 to 21~~, wherein R⁷ represents a group having an electron withdrawing inductive effect.

23. (Original) Compound according to claim 22, wherein R⁷ is selected from the group consisting of: halogen, C₁-C₄ alkoxy.

24. (Original) Compound according to claim 23, wherein R⁷ represents a halogen.

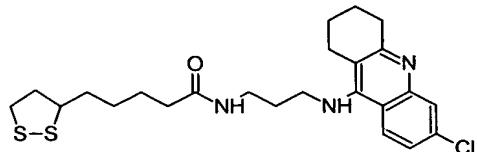
25. (Currently amended) Compound according to ~~one of the claims from~~ claim 17 ~~to 24~~, wherein R⁷ is selected from the group consisting of: halogen, hydrogen, methoxy; R⁵ is selected from the group consisting of: hydrogen, amine, nitroalkyl, halogen, hydroxy.

26. (Currently amended) Compound according to ~~one of the claims from~~ claim 17 to 25, wherein R⁷ is situated in position 6.

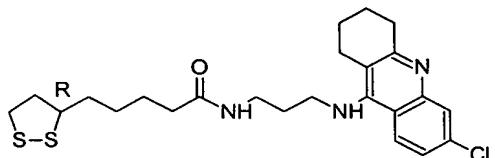
27. (Currently amended) Compound according to ~~one of the claims from~~ claim 17 a 26, wherein R⁵ is selected from the group consisting of: hydrogen, C₁-C₄ amine, C₁-C₄ nitroalkyl, -NH₂, nitro, halogen.

28. (Currently amended) Compound according to ~~one of the claims from~~ claim 17 to 26, wherein R⁵ is selected from the group consisting of: hydrogen, halogen.

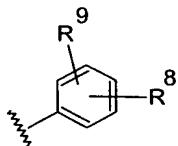
29. (Original) Compound according to claim 28, and having the following formula:



30. (Original) Compound according to claim 29, in form R:



31. (Original) Compound according to claim 6, wherein Ar presents the formula:



wherein R¹ represents a C₃-C₉ alkandiamine.

32. (Original) Compound according to claim 31, wherein R¹ represents a C₆-C₈ alkandiamine.

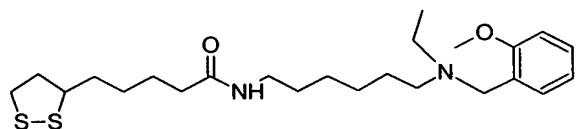
33. (Currently amended) Compound according to claim 31 or 32, wherein R¹ presents the formula -NR¹⁶-R¹⁷-NR¹⁸-R¹⁹-, wherein R¹⁹ is linked to Ar and -NR¹⁶ is linked to the carbonylic carbon; R¹⁷ is a C₂-C₇ alkyl; R¹⁶ and R¹⁸ are selected, each independently of the other, from the group consisting of: C₁-C₃ alkyl, hydrogen; R¹⁹ represents a C₁-C₃ alkyl.

34. (Original) Compound according to claim 33, wherein R¹⁷ is a C₃-C₆ alkyl; R¹⁶ represents a hydrogen; R¹⁸ is selected from the group consisting of: ethyl, methyl, hydrogen; R¹⁹ represents a methyl.

35. (Currently amended) Compound according to ~~one of the claims from~~ claim 31 or 34, wherein R⁹ is selected from the group consisting of: hydrogen, hydroxy, halogen, C₁-C₄ alkoxy; R⁸ is selected from the group: hydroxy, halogen, C₁-C₄ alkoxy.

36. (Original) Compound according to claim 35, wherein R⁹ represents a hydrogen and R⁸ represents a methoxy situated in ortho or meta position with respect to the remaining part of the compound.

37. (Original) Compound according to claim 36, and having the following formula:



38. (Currently amended) Compound having the general formula (I), as defined in ~~any one of the claims from claim 1 to 37~~, for use as a medicament.

39. (Cancelled)

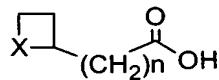
40. (Cancelled)

41. (Cancelled)

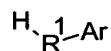
42. (Currently amended) Pharmaceutical preparation comprising a compound having general formula (I), as defined in ~~any one of the claims from claim 1 to 37~~, or a pharmaceutically acceptable salt, and an excipient and/or pharmaceutically acceptable diluent.

43. (Currently amended) Method for the treatment of Alzheimer's disease in a mammal; ~~the method comprises comprising~~ administering to said mammal an efficacious quantity of a compound having general formula (I), as defined in ~~any one of the claims from claim 1 to 37~~.

44. (Currently amended) Method for the synthesis of a compound having general formula (I), as defined in ~~any one of the claims from claim 1 to 37~~, comprising an addition phase wherein a compound having the general formula (II):



is reacted with a compound having the general formula (III):



in basic conditions.

45. (New) Method for the treatment of a pathology characterized by deposits of β -amiloid ($\text{A}\beta$) in mammals comprising administering to said mammal an efficacious quantity of a compound having a general formula (I), as defined in claim 1.